

# EXHIBIT 1

UNITED STATES DISTRICT COURT

DISTRICT OF MINNESOTA

In Re: Bair Hugger

Forced Air Warming

Products Liability

Litigation

This document relates ) MDL No. 15-2666

to all actions )

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The videotaped deposition of JIM HO, in the  
above-styled suit, was taken pursuant to notice for  
discovery and/or evidentiary purposes, before Donna  
Gerbrandt CSR(A), at the offices of Borden Ladner  
Gervais LLP, Calgary, Alberta, Canada, on the 28th  
day of June, 2017.

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1 (Proceedings commenced at 8:14 a.m.)

2 THE VIDEOGRAPHER: Here begins the  
3 videotaped deposition of Jim Ho in the matter of  
4 Re: Bair Hugger forced air warming products  
5 liability litigation, in the United States District  
6 Court, District of Minnesota, MPL No. 15-2666.

7 Today's date is June 28th, 2017.

8 The time on the video monitor is 8:15 a.m. We are  
9 on the record.

10 The videographer today is  
11 Bernice Dubon on behalf of Amicus Reporting Group.  
12 The video deposition is taking place at the offices  
13 of Borden Ladner Gervais of Calgary, Alberta.

14 Would counsel please voice identify  
15 themselves and state whom they represent.

16 MR. BANKSTON: Mark Bankston on  
17 behalf of the plaintiffs.

18 MR. ASSAAD: Gabriel Assaad on  
19 behalf of the plaintiffs.

20 MR. GORDON: Corey Gordon on  
21 behalf of the defendants 3M and Arizant.

22 THE VIDEOGRAPHER: The court reporter  
23 today is Donna Gerbrandt on behalf of  
24 Amicus Reporting Group. Would the reporter please  
25 swear in the witness.

1 JIM HO, sworn

2 BY MR. BANKSTON

3 Q. Good morning, Mr. Ho.

4 A. Good morning.

5 Q. I'm going to be talking to you today  
6 about some opinions that you gave in this case. I  
7 understand -- you've given a deposition before;  
8 right?

9 A. That's true.

10 Q. Okay. How many times do you think  
11 you've been deposed before?

12 A. Once.

13 Q. One time. Okay. Then just as a little  
14 refresher, I know you don't do this all the time,  
15 it's just like we're in a courtroom, just there's  
16 no judge here. We're going to be asking each  
17 other -- I'm going to be asking you questions,  
18 you'll be giving me answers. We need to be really  
19 careful not to talk over each other. She's writing  
20 everything down. So we'll try to pause between  
21 each other. You know, sometimes in natural  
22 conversation you tend to interrupt each other,  
23 finish each other's sentences. It's tough to do  
24 that for her so we'll try to avoid that.

25 I also know, not being an

1           your education. I know you have a bachelors and a  
2           masters in microbiology?

3           A.    True.

4           Q.    Okay. The PhD is in microbial  
5           chemistry?

6           A.    Right.

7           Q.    Do you have any other advanced  
8           education?

9           A.    Yes. I spent a lot of time  
10          understanding aerosol technology. The nature of  
11          the work I was doing required that I understand  
12          all -- all the subjects involved with aerosol  
13          technology.

14          Q.    Okay. Is that -- is that knowledge you  
15          acquired at the Department of National Defence?

16          A.    It's mostly through attaining --  
17          attending conferences --

18          Q.    Okay.

19          A.    -- working with colleagues, and mostly  
20          hands-on, intending to use instrumentation to  
21          characterize and define aerosol characteristics.

22          Q.    Okay. You would agree that your career  
23          has been spent as a researcher for the Canada's  
24          Department of National Defence?

25          A.    That's correct.

1 Q. In your report you divided your career  
2 up into pre 1990s and post 1990s. Is that a good  
3 way to divide your career, you think?

4 A. Yeah, yes.

5 Q. Okay. I understand that the first part  
6 of your career was what we were just talking about,  
7 correct, the understanding of biological aerosols?

8 A. Yeah.

9 Q. Okay. And you would agree an aerosol is  
10 either a solid or a liquid particle suspended in  
11 the air?

12 A. Say again.

13 Q. An aerosol is a solid or a liquid  
14 particle suspended in the air?

15 A. That's right.

16 Q. Okay. You would agree that aerosols  
17 travel along the current of the air?

18 A. True.

19 Q. Okay. Now, your career post 1990s has  
20 been primarily of the development of biological  
21 detection systems; correct?

22 A. Yes, but the -- but the mandate from the  
23 very beginning of my career was to develop  
24 biological detection technologies.

25 Q. Okay.

1           developing the technology.

2           Q.    Okay.  And would you agree that for  
3           nearly 30 years you've been in the practice of  
4           developing devices and using -- using or selling  
5           them to various clients and the government and  
6           others?

7           A.    That's not exactly true.  My job is to  
8           develop the hardware and the software technology  
9           for military applications, which is Canadian  
10          military, and -- and selling it really was not my  
11          job.

12          Q.    Okay.  What hardware and software  
13          technologies did you use in this case?

14          A.    Well, the final instrument that  
15          performed the task of detecting the presence of  
16          live agents was what we would call -- it's an  
17          abbreviation, F-L-A-P-S, FLAPS.

18          Q.    Okay.

19          A.    So I found that it is not just good  
20          enough to develop an instrument that you think is  
21          good enough.  You actually have to go out --  
22          outside of the laboratory and demonstrate that it  
23          works in the -- in the environment it was -- it was  
24          set for, which is military -- military conditions,  
25          which is outdoors.

1 to determine what the name of the microorganism is.  
2 And then once you discover that, then you know  
3 whether it is going to be disease causing or not.

4 Does that go along with what you're  
5 expecting?

6 Q. Not exactly, but let's get more  
7 specific. And I think that will help us, if we get  
8 a little more specific. For instance, are you an  
9 expert in evaluating clinical outcomes in  
10 healthcare settings?

11 A. No.

12 Q. Okay.

13 A. Yeah.

14 Q. Are you an expert in hospital air  
15 quality?

16 MR. GORDON: I object to the form  
17 of the question.

18 A. I -- I'm not sure what you're trying to  
19 understand from that question, but you would notice  
20 that we have done some experiments with the -- with  
21 the wind tunnel to determine how you may be able to  
22 detect presence of biological particles in a clean  
23 room condition. So if you could -- if you could  
24 transpose that into what you're asking, then the  
25 answer would be yes. If you don't accept that,



1 jury of 12 Americans that you're an expert in the  
2 field of microbiology; correct?

3 A. I -- I would say that if that's what  
4 they consider having spent 30 years in -- in bio  
5 aerosol as an expert, then that is fair enough.

6 Q. Okay. Are you an expert in operating  
7 rooms?

8 A. No.

9 Q. Okay.

10 A. No.

11 Q. Are you -- do you have any expertise on  
12 the levels of bioburden within an operating room?

13 A. No.

14 Q. Okay. So you wouldn't be able to tell  
15 me, for instance, does every area in an operating  
16 room have equivalent levels of bioburden?

17 A. I -- I cannot say with a blanket  
18 statement.

19 Q. Okay.

20 A. Nor can anybody else really.

21 Q. Well, would you agree with me -- do you  
22 have any expertise to say whether the area  
23 underneath the surgical table has more bioburden  
24 than other parts of the room?

25 A. No. Yeah.

1 Q. You're not an expert in orthopedic  
2 surgery, are you?

3 A. No. Apart from the fact that I'm the  
4 son of an orthopedic surgeon. Does that help any?

5 Q. Again, I'm not -- I'm not answering  
6 questions today.

7 A. Yeah.

8 Q. I can't -- I can't help you along. You  
9 know, that's -- I can let you answer questions is  
10 what I can do.

11 A. Yeah. Yeah.

12 Q. But do you -- do you think that you  
13 could comfortably represent to a jury of 12  
14 Americans that you being the son of an orthopedic  
15 surgeon makes you an expert qualified to give  
16 opinions in a lawsuit?

17 A. That would be a bit far fetched.

18 Q. I would think so. Okay. You're not an  
19 expert in anesthesiology, are you?

20 A. No.

21 Q. You're not an expert in infectious  
22 disease?

23 A. It would be safe to say no, but in the  
24 work area that I'm in I have to be aware of what  
25 are the threat agents for the Canadian military,

1 and for that matter we need to know a lot about  
2 infectivity, if that's what you're driving at.

3 Q. Can you describe to me what a  
4 peri-prosthetic joint infection is?

5 A. No.

6 Q. Okay. So do you have any -- I mean, I  
7 assume since not giving a definition, you wouldn't  
8 consider yourself as having specific expertise in  
9 peri-prosthetic joint infections?

10 A. No.

11 Q. Okay. Did you know anything about the  
12 device before accepting work in this case? And by  
13 "the device" I mean the Bair Hugger surgical  
14 warming unit.

15 A. No.

16 Q. You're not an engineer; correct?

17 A. That's correct.

18 Q. You're not a biomedical engineer?

19 A. No.

20 Q. What do you understand the device to do,  
21 the Bair Hugger?

22 A. From the description that was -- that I  
23 read and what was told to me, it's a technology or  
24 a device that provides warming features for a  
25 patient under -- under operation.

1 Q. Okay. Have you ever seen the device?

2 A. No.

3 Q. You would agree with me that as an  
4 expert you need to understand the nature of the  
5 problem being claimed in order to investigate it?

6 MR. GORDON: I object to the form  
7 of the question.

8 A. I -- I need to -- to know exactly  
9 what -- what is the -- the issues at hand.

10 BY MR. BANKSTON

11 Q. Yeah, that's a good way to put it.  
12 Like, for instance, in a lawsuit the issues at hand  
13 is what the plaintiff is claiming happened to him  
14 because of this device; right?

15 MR. GORDON: I object to the form  
16 of the question, also lack of foundation.

17 BY MR. BANKSTON

18 Q. Would you agree with that?

19 A. Well, if I -- if I were to give -- give  
20 an impression of what biological aerosols are,  
21 would I need to know any of the things that you say  
22 I need to know?

23 Q. Well, I haven't even said anything. I'm  
24 wondering what do you think the issue is that  
25 you're here to address?

1           Q.    Okay.  For instance, do you understand  
2           what the plaintiffs are claiming happened to them  
3           in this case?

4           A.    If I understand correctly, they claim  
5           that the air coming from a Bair Hugger has the  
6           potential to cause infections.

7           Q.    You mean -- and you mean the air being  
8           exhausted out of the Bair Hugger?

9                   MR. GORDON:           I object to the form  
10          of the question.

11          A.    I -- I -- I wouldn't know if it is  
12          exhausted or coming from it, but that's the overall  
13          impression.

14                   BY MR. BANKSTON

15          Q.    Have you seen the plaintiffs' complaint  
16          before?

17          A.    What do you mean?

18          Q.    Oh, okay.  I forget.  I'm asking you  
19          questions like you're an expert who comes to  
20          depositions every week.  When I say "complaint" --

21          A.    Yeah.

22          Q.    -- actually that's a legal term --

23          A.    Yeah.

24          Q.    -- meaning the initiating document of a  
25          lawsuit.  Have you ever seen that document where

1 the plaintiff sets forth why they think 3M is  
2 responsible for something?

3 A. I don't believe so.

4 Q. Okay. Do you have an understanding of  
5 how the plaintiffs believe the Bair Hugger caused  
6 their infections?

7 A. I don't -- I don't think so.

8 Q. Okay. The first topic I really want to  
9 talk to you about in your report is regarding the  
10 size of particles. You know that there's a  
11 discussion about the size of particles and the size  
12 of biological aerosols?

13 A. Right.

14 Q. Okay. Did you bring a copy of your  
15 report with you today?

16 A. I did.

17 Q. Okay. Do you want to get that out for  
18 me and we'll take a look at it together.

19 Now, Mr. Ho, before we dive into  
20 the report itself, I want to talk generally in  
21 terms of what you reviewed to create this report.  
22 From what I understand, everything that you  
23 reviewed is cited somewhere in the report?

24 A. Yeah. It's open literature material.

25 Q. Okay. So you would agree with me that

1 A. I don't think I say that.

2 Q. Okay. You gave opinions about filters;  
3 right?

4 A. Yeah.

5 Q. Okay. You know what I mean when I talk  
6 about a MERV 14 filter?

7 A. Yes.

8 Q. Okay. You believe that a MERV 14 filter  
9 is adequate inside the Bair Hugger; correct?

10 A. Yes.

11 Q. You believe that HEPA filters are  
12 overkill for this application?

13 A. That is a quotation from a source.

14 Q. Do you believe that HEPA filters are  
15 overkill in this application?

16 A. Are you wanting an opinion right here  
17 now?

18 Q. Hmm hmm. That's what you're here for,  
19 sir.

20 A. Yeah, I think HEPA filters are more than  
21 what would be required.

22 Q. Required for what?

23 A. For the -- the purpose of what the  
24 instrument is supposed to do.

25 Q. Okay. And in terms of the filter, what

1 correct?

2 A. That's right.

3 Q. Okay. You will agree with me, though,  
4 that the reason you want to keep particles out of  
5 an operating room to an absolute minimum is to  
6 prevent the incident of surgical infection;  
7 correct?

8 A. Now, where are we going with this one  
9 again? You already said that I'm not -- I'm not an  
10 expert in clean-room facilities.

11 Q. Okay.

12 A. And why are you asking me that question  
13 again?

14 Q. So you can tell me that's not a question  
15 you're qualified to answer?

16 A. Yeah.

17 Q. Okay.

18 A. I'm simply -- I'm simply here to provide  
19 you with insight into -- into bio aerosol  
20 technologies.

21 Q. Okay. And I appreciate that. I want  
22 you to tell me whenever that's true.

23 A. Yeah. Yeah.

24 Q. Whenever I'm talking or asking you a  
25 question about something you're not qualified to



1 talk about, tell me "That's not why I'm here,  
2 Mr. Bankston. I'm here for a totally different  
3 reason."

4 A. Yeah.

5 Q. That's totally fine. I don't have any  
6 problems with that. Let's talk a little bit more  
7 about this MERV 14 filter. You say -- let's go to  
8 page 25 of your report.

9 A. Got it.

10 Q. Okay.

11 MR. GORDON: Did you say 24?

12 MR. BANKSTON: 25.

13 Q. All right. Do you see the section that  
14 starts with D. --

15 A. Hmm hmm.

16 Q. -- MERV 14 filtration?

17 A. Yeah.

18 Q. Okay. You see the second sentence in  
19 that paragraph?

20 A. Yeah.

21 Q. It says: "Standard charts list this  
22 specification: removal of all bacterial particles  
23 sized within .3 to 1 micron."

24 A. Yeah.

25 Q. Do you see that?

1 Q. Yes. Yes. So you understand that this  
2 purpose is to provide an assessment of the filter  
3 efficiency on the Bair Hugger system and the  
4 efficiency level of the current filters? Do you  
5 see that's what this document says?

6 A. Right.

7 Q. And you see how there is a  
8 "3M Confidential" up at the top of the document?

9 A. Hmm hmm.

10 Q. Okay. On the bottom of -- the bottom  
11 corner of this document you'll see a number that  
12 starts with 3MBH; correct?

13 A. Right.

14 Q. Okay. And the number here starts with  
15 89. The part of this document that I would like to  
16 ask you about today is going to be -- the final  
17 numbers are going to be 96. So if you can flip to  
18 96 for me. That's the part I would like to ask you  
19 a question about.

20 Okay. Perfect. Now, you see that  
21 there's been a portion of that document that's been  
22 highlighted that reads "Table 3. No load (initial)  
23 tests for Model 775 filter." Do you see where that  
24 is?

25 A. Yeah.

1           this table, the Bair Hugger filter does not remove  
2           all particle sizes within .3 to 1 micron?

3           A.     That's what it says.

4           Q.     Correct. In fact you would also agree  
5           with me that the Bair Hugger filter does not even  
6           remove all particles between 1 and 3 microns;  
7           correct?

8           A.     Right.

9           Q.     Now, that's a little bit different than  
10          the standard specification that you discussed in  
11          your report; correct?

12                   MR. GORDON:           I object to the form  
13          of the question. It misstates the evidence.

14          A.     Are you referring to the 99 percent  
15          numbers?

16                   BY MR. BANKSTON

17          Q.     No. What I'm actually referring to is  
18          remember when you told me that a MERV 14 filter by  
19          specification will remove all particles sized .3 to  
20          1 micron? Do you remember telling me that?

21          A.     Right.

22          Q.     Now, that is not -- this Bair Hugger  
23          filter test that you have in front of you, that  
24          does not meet that standard, does it?

25          A.     It does appear to be slightly different.

1           Q.    In other words, from this chart we can  
2           see -- let's go down all four lots that were  
3           tested. In the first lot it only removed  
4           83 percent of those particles; correct?

5           A.    Yeah.

6           Q.    In the second test it only removed  
7           82 percent; correct?

8           A.    Yeah.

9           Q.    In the next test it only removed  
10          75 percent; correct?

11          A.    Yeah.

12          Q.    And in the next test it only removed  
13          78 percent; correct?

14          A.    Correct.

15          Q.    So it does not meet the standard in  
16          which you expressed in your report; correct?

17          A.    Right.

18          Q.    Okay. Thanks. Are you familiar with  
19          what a HEPA filter is?

20          A.    Yes.

21          Q.    Okay. Are you familiar with the  
22          specifications for a HEPA filter?

23          A.    I don't have the numbers handy, but in  
24          general.

25          Q.    Let me throw out a number and see if it

1 MR. GORDON: I object to the form  
2 of the question. That is not -- you misread it.

3 BY MR. BANKSTON

4 Q. All right. Let's read the whole thing.  
5 "According to table 8.2 of Kowalski, a MERV 14  
6 filter will remove Staph. aureus with  
7 97% efficiency..." Is that a correct reading of  
8 that?

9 A. Yeah.

10 Q. So by some fairly simple subtraction we  
11 know that 3 percent of staph aureus organisms will  
12 pass through this filter; correct?

13 A. You can assume that.

14 Q. Okay. Do you agree that in selecting a  
15 filter for use in a healthcare setting you need to  
16 know the environment in which it's going to be  
17 used?

18 MR. GORDON: I object to the form  
19 of the question. Vague, ambiguous, lack of  
20 foundation, incomplete hypothetical.

21 A. There is a "but" to that question?

22 BY MR. BANKSTON

23 Q. A "but?"

24 A. Yeah. Do you have some follow-up to  
25 that question?

1 Q. I'm sure I'll have more questions, yeah.

2 A. Yeah. So what is the question again?

3 Q. When you're selecting a filter for use  
4 in a healthcare setting, do you need to know the  
5 environment of use?

6 MR. GORDON: I object to the form  
7 of the question.

8 A. If I were designing an instrument? Is  
9 that what you're saying.

10 BY MR. BANKSTON

11 Q. No. I'm actually asking if you're  
12 selecting a type of filter for use in a healthcare  
13 setting. Not if you're making a device. Like just  
14 if you're picking a filter. If you're going to  
15 pick a filter, do you need to know the environment  
16 it's going to be used in?

17 A. That's sort of a vague question, though,  
18 because it's hard to really answer that question  
19 unless I know what is it that you really want to  
20 point at.

21 Q. Okay.

22 A. There seems to be a second part to that  
23 question, depending on whether the answer is yes or  
24 no.

25 Q. Okay. So in terms of --

1           A.    Come right out to the question and  
2           see --

3           Q.    That's my question.  I'm wondering --  
4           let's say I have a job, and my job is to pick a  
5           filter.

6           A.    Pick a filter.

7           Q.    I'm going to pick a filter for an  
8           application.

9           A.    Yeah.

10          Q.    Do I need to know where the filter is  
11          going to be used if I'm going to pick that filter  
12          safely?

13          A.    It would be helpful.

14          Q.    In order to determine if a filter is  
15          safe in a given application, you might need to know  
16          the environment of use; correct?

17          A.    When you say "safe," how would -- how  
18          would that mean?  Is that an absolute term or is it  
19          a --

20          Q.    That's a good point.

21          A.    -- is it something that is adequate for  
22          the job?

23          Q.    Yeah.  Let's phrase it in the way it's  
24          done in your report, for instance.  You say that a  
25          MERV 14 filter is adequate for this application?

1 A. Yeah.

2 Q. I would assume when you speak of  
3 "adequate," that means reasonable in terms of  
4 patient safety as well; right?

5 A. Yeah. That's a bit of a stretch,  
6 though.

7 Q. So let me make sure I have this clear.  
8 When you say that a MERV 14 filter is adequate,  
9 you're not talking about patient safety?

10 A. You -- you really want to narrow it down  
11 to what -- what the issue is at hand.

12 Q. That's absolutely why we're here, yeah.

13 A. Yeah.

14 Q. Right. Okay. So let me ask it again.

15 A. Right.

16 Q. Okay. When you say in your report, your  
17 words, a MERV 14 filter is adequate in this  
18 Bair Hugger --

19 A. Yeah.

20 Q. -- do you mean from a patient safety  
21 point of view?

22 A. A patient safety could be interpreted in  
23 a variety of directions. So in this case you  
24 are -- you're really trying to equate filter X,  
25 safety, yes; filter Y, safety no, and I can't



1 answer that question.

2 Q. Okay. So you cannot give the opinion  
3 that the Bair Hugger filter is adequate from a  
4 patient safety perspective?

5 A. I -- I can say that the filter selected  
6 is adequate for the performance of the instrument.  
7 And I want to take safety out of it because --  
8 because safety is a whole different issue.

9 Q. Okay. So in terms of -- you're a  
10 designer of devices; correct?

11 A. I do some of that, yeah.

12 Q. Okay. And so sometimes when making a  
13 device you have to understand if a component that  
14 you're using in the device is going to have a  
15 negative effect and make your device not work or  
16 whether it will work fine with that component. Is  
17 that simple enough?

18 A. Hmm hmm.

19 Q. Okay. So what I -- oh, is that a yes?

20 A. Yes.

21 Q. And I don't mean to be rude about it.

22 A. Yeah. Yeah.

23 Q. She can't take down --

24 A. Right. Right. Sorry.

25 Q. No problem. So am I correct, when you

1 say the Bair Hugger filter is adequate, that's in  
2 terms of the function of the device?

3 A. Yes.

4 Q. Okay.

5 A. Yeah.

6 Q. So what I want to make sure, so when one  
7 day if we get to trial, is you're not making any  
8 representations to this jury about whether that  
9 Bair Hugger filter is adequate from a patient  
10 safety standpoint?

11 A. Again I like to emphasize the fact that  
12 when you -- when you attach safety and -- and  
13 selection of material, then you are making a very  
14 huge leap in faith in saying that. So -- so I'm  
15 only speaking from the standpoint of an aerobiology  
16 technical person. So the future that is selected  
17 is adequate for what the instrument is supposed to  
18 do.

19 Q. You mean it's adequate for the device to  
20 be able to blow hot air on the patient to warm them  
21 for surgery?

22 A. It's adequate to provide the airflow  
23 characteristics that -- that the end result is  
24 called for.

25 Q. Okay. In terms of is that filter

1 sufficient to provide reasonable assurance that a  
2 patient will not suffer peri-prosthetic joint  
3 infection, that's probably not something you can  
4 talk about today?

5 A. No.

6 Q. Okay. Let me ask that in another way  
7 because I want to be very specific, not just about  
8 peri-prosthetic joint infection. You would agree  
9 with me you do not have the necessary  
10 qualifications and expertise to state the level of  
11 filtration that is needed in the Bair Hugger to  
12 maintain clinically safe levels of air quality in  
13 an ultra clean operating room during an orthopedic  
14 procedure?

15 MR. GORDON: I object to the form  
16 of the question.

17 BY MR. BANKSTON

18 Q. Do you agree with that? And if you need  
19 to, I can repeat it and we can do it again.

20 A. You're saying if I do or do not have an  
21 expertise in designing something?

22 Q. No, no, no. This is very, very specific  
23 so let's go one -- let's go real slow.

24 A. Okay.

25 Q. Okay. What I'm asking is are you

1           that Elghobashi relied upon and that counsel was  
2           incorrect.

3                       MR. BANKSTON:           I thought it was  
4           Kalliomaki, but I guess not. I guess he could  
5           have.

6                       MR. ASSAAD:           No. No. Elghobashi  
7           went in a different.

8                       MR. GORDON:           Yeah, Villafruela.

9                       MR. BANKSTON:           Yeah. It was an  
10          isolation room, yeah, but performed before an  
11          operating room. We'll still get some mileage out  
12          of Kalliomaki, though. Don't worry about that.

13                      BY MR. BANKSTON

14                      Q. All right. Let's talk a little bit  
15          about particle count. You're familiar with the  
16          practice of particle counting; right.

17                      A. Yes.

18                      Q. You don't believe that particle counting  
19          can be predictive of microbiological contamination  
20          of air in an operating room, do you?

21                      A. I do not.

22                      Q. Okay. A lot of people disagree with you  
23          about that. Do you recognize that?

24                      MR. GORDON:           I object to the form  
25          of the question, lack of foundation, assumes facts

1                   Now, you would agree with me that  
2           the conclusion of Mr. Stocks -- excuse me, of  
3           Dr. Stocks and his team is that particles are a  
4           reasonable surrogate for bioburden?

5           A.    Why do I have to agree with you?

6           Q.    You don't have to.  I'm asking if you  
7           do.

8           A.    Yeah.  The answer is no.

9           Q.    No, you don't agree with that.  So if  
10          somebody was to say Stocks and his colleagues were  
11          able to demonstrate that particles are a reasonable  
12          surrogate for bioburden, you would say no, that's a  
13          wrong opinion?

14          A.    Correct.

15          Q.    Okay.  Are you familiar with a  
16          Russell Olmsted with the National Institute of  
17          Health?

18          A.    What is this in relation to?

19          Q.    I'm just wondering if you know the man.

20          A.    No.

21          Q.    Okay.

22          A.    Yeah.

23                   MR. BANKSTON:           4.

24          Q.    Okay.  So one of the things we had  
25          talked about with Stocks, right, is the people who

1 have studied this, is they haven't really followed  
2 the right methodology to really measure this. Is  
3 that part of your contention?

4 MR. GORDON: Objection, form of  
5 the question.

6 A. If you were to look at one of the figure  
7 studies he's presented --

8 BY MR. BANKSTON

9 Q. Hmm hmm.

10 A. His Figure 1. Do you see it?

11 Q. Sure. Yeah. Let's look at Figure 1.

12 A. Yeah.

13 Q. Okay.

14 A. Okay. He's based all his conclusions on  
15 Figure 1.

16 Q. Okay. So --

17 A. And if you -- if you have any training  
18 in measurements, particle analysis, and all the  
19 other good things that one would have to have, you  
20 would look at that Figure 1, right away would say  
21 that this is scattered data all over the map.

22 Q. Okay.

23 A. Wouldn't you agree?

24 Q. I don't have any expertise to -- I would  
25 rely on somebody who is an expert.

1           A.    Okay.  So as a person who has measured  
2           particles, live agent, everything else, that's  
3           definitely bad data.

4           Q.    Anybody who has measured particles for a  
5           living would know that's bad data?

6           A.    Yeah.

7           Q.    Okay.

8           A.    Very bad.

9           Q.    Very bad?  Okay.

10          A.    And his whole conclusion is based on  
11          that observation.

12          Q.    Okay.  So there's, according to you,  
13          some methodological problems and this is not good  
14          stuff?

15          A.    On top of that the data was fudged.

16          Q.    Okay.  The data was fudged?

17          A.    Yeah.

18          Q.    What do you mean by that?

19          A.    Well, somewhere along the way his raw  
20          data did not fit his expectations so he took it  
21          upon himself to do a data transform.

22          Q.    Why do you say that?  Where do you get  
23          that from?

24          A.    He said that.

25          Q.    Okay.  Where is that in his -- oh,

1 BY MR. BANKSTON

2 Q. Now let's move on.

3 MR. GORDON: It misstates the  
4 evidence.

5 BY MR. BANKSTON

6 Q. The second -- the second paragraph of  
7 that email.

8 A. Yeah.

9 Q. Do you have Exhibit 26 in front of you?

10 A. Yeah.

11 Q. Now, Mr. Olmsted says that he has done  
12 investigations where he used electronic particle  
13 counts; correct?

14 A. He said that, yeah.

15 Q. Okay. And then he says: "...it appears  
16 this group was able to demonstrate particle counts  
17 serve as a reasonable surrogate for bioburden of  
18 air in an OR."

19 A. Yeah.

20 Q. You disagree with that?

21 A. Totally.

22 Q. Okay. And I believe you also told me  
23 that anybody who has done particle counting would  
24 immediately recognize that that's not true?

25 A. Correct.



1 correlates with airborne colonies  
2 and represents an acceptable  
3 surrogate for daily assessment of  
4 cell-processing cleanroom  
5 performance"

6 A. So are we done with the Stocks paper?

7 BY MR. BANKSTON

8 Q. Yeah, we're done with that. You can put  
9 that away.

10 I've handed you, sir, what has been  
11 marked for the purposes of this deposition as Ho  
12 Exhibit 1. Do you see in front of you a paper by  
13 Raval et al, and I'm going to read the title.  
14 "Real-time monitoring of non-viable airborne  
15 particles correlates with airborne colonies and  
16 represents an acceptable surrogate for daily  
17 assessment of cell-processing cleanroom  
18 performance." Did I read that title correctly?

19 A. You read the title correctly.

20 Q. This is not something that you reviewed  
21 in coming to your opinions in this case; correct?

22 A. That's correct.

23 Q. Have you ever seen this study before?

24 A. No.

25 Q. Okay. Do you see on the results where

1           it says "viable and nonviable particles were well  
2           correlated?"

3           A.    It says that.

4           Q.    Describe what it means to me if those  
5           particles are well correlated? What does "well  
6           correlated" mean?

7           MR. GORDON:           Well, you're going  
8           to have to give him the opportunity to read the  
9           study if you want him to comment on specifics.

10          BY MR. BANKSTON

11          Q.    Well, I just want to know what  
12          "correlation" is. Do you know what it means when  
13          things are correlated, two different findings are  
14          correlated?

15          A.    Let me -- can I flip through some of the  
16          results and interpretation --

17          Q.    Yeah. I'm not going to stop you.

18          A.    -- before I -- before I get too deep  
19          into this thing?

20          Q.    Hmm hmm. Dr. Ho, if you just want to  
21          read the whole thing, I'm going to take a little --

22          A.    Yeah, sure.

23          MR. BANKSTON:           Go off the record  
24          for a second.

25          THE VIDEOGRAPHER:    We are going off the

1 A. Yeah. Yeah.

2 Q. So when it comes to whether forced air  
3 warmers affect the risk of surgical site infection  
4 in operating rooms, you don't have an opinion?

5 A. No.

6 Q. Thank you, sir.

7 A. Yeah.

8 Q. The next sentence states:

9 "Other areas of the hospital  
10 caring --"

11 Excuse me.

12 "Other areas of the hospital caring  
13 for high-risk patients with  
14 increased risk of nosocomial  
15 infection, such as burn units and  
16 hematology/oncology wards, have put  
17 air monitoring and quality systems  
18 into place..."

19 Do you see that?

20 A. I see that.

21 Q. Okay. The final sentence states:

22 "Thus reduced airborne particulates  
23 appear to correlate with a  
24 decreased risk of nosocomial  
25 infections in high-risk patient

1                   populations."

2                   Do you have any opinions about whether  
3                   that statement is scientifically valid or not?

4                   A.    That statement came out of the blue.  It  
5                   has got no real backing to it.

6                   Q.    Okay.  So your opinion is that statement  
7                   has no support and is not true?

8                   A.    No.

9                   Q.    Okay.  Yes?  I'm sorry, it's --

10                  A.    No.  No.  No.

11                  Q.    And the question asked is you said it  
12                  was not true.  There's the negative thing, and I  
13                  think you're saying the opposite of what the  
14                  transcript is going to reflect.

15                  A.    It's not -- it's not true.

16                  Q.    So you're saying this statement is not  
17                  true?

18                  A.    Yeah.

19                  Q.    So this is another piece of  
20                  peer-reviewed literature which disagrees with you,  
21                  which you say is wrong?

22                  A.    Correct.

23                  Q.    Okay.  These -- these authors here, I  
24                  take it you also say that they don't know what  
25                  they're doing either; right?

1 Q. Okay. So this is not something that you  
2 relied on when coming to your opinions about  
3 whether particles are proxies for bioburden?

4 A. Come again.

5 Q. This is not something that you reviewed  
6 when coming to your opinion that particles are not  
7 proxies for bioburden?

8 A. Yeah, I haven't seen this paper before.

9 Q. Okay. I'll tell you what? You want to  
10 take -- this one isn't too long, actually. This  
11 one only has about -- it looks like eight pages of  
12 text. If you want to review this to see if there's  
13 anything you want to look at before I start asking  
14 you questions, I'm going to ask you a few questions  
15 about this before we go to lunch.

16 A. Okay.

17 MR. GORDON: Did you say  
18 Darouiche was a microbiologist?

19 MR. BANKSTON: No. That him being  
20 in microbiology, I thought he might be familiar  
21 with his work in microbiology. Because, as you  
22 see, it's a study of airborne microorganisms. I  
23 thought it might have hit his Google alerts.

24 A. Do you have a copy that is not smeared?

25 Q. Yeah. That I sure don't, unfortunately.

1 Q. In this lawsuit --

2 A. Yeah.

3 Q. -- what you're here to testify about --

4 A. Right.

5 Q. -- is implant infections?

6 A. Right.

7 Q. Okay. You understand that the  
8 mechanisms, the biological, the physiological  
9 mechanisms by which an infection happens  
10 incisionally versus peri-prosthetically are  
11 different mechanisms? Do you have enough expertise  
12 to know that?

13 A. I don't --

14 MR. GORDON: I object to the form  
15 of the question.

16 A. Yeah. I think it's a bit technical for  
17 me here.

18 BY MR. BANKSTON

19 Q. Okay. Let's move on then to -- into  
20 that paragraph. That first part we were talking  
21 about was one of their findings. Another finding  
22 that they had was -- you see right after it says  
23 "Figure 4" there's a new sentence. And it says:

24 "CFU density was positively related  
25 to total particulate density ... in

1           the control group, indicating that  
2           airborne [particulate] counts may  
3           be used as a proxy for ambient CFU  
4           density."

5           I want to ask you about some terms that  
6           are used in there. First, because I don't think  
7           we've defined it so far, a CFU is a colony-forming  
8           unit; correct?

9           A. Yeah.

10          Q. Okay.

11          A. Yeah.

12          Q. And in this case, when they talk about  
13          CFU density, we can think about that as the amount  
14          of -- the total amount and concentration of  
15          airborne biological mass that they're measuring;  
16          correct?

17          A. Right.

18          Q. Okay. And their conclusion is, is that  
19          airborne particle counts correlate well and can be  
20          used as a proxy for the CFUs.

21          A. What do you make of the following  
22          sentence after that?

23          Q. We're going to keep going. Lets stay  
24          one step at a time.

25          A. Yeah.

1 Q. Stop trying to figure out where we're  
2 going here, Mr. Ho. Let's try to answer the  
3 questions that are in front of you.

4 You understand that they found that  
5 airborne particle counts may be used as a proxy for  
6 ambient CFU density? That's what their statement  
7 there says?

8 MR. GORDON: I object to the form  
9 of the question.

10 BY MR. BANKSTON

11 Q. Correct?

12 A. That's what they say here.

13 Q. And that's something that you disagree  
14 with?

15 A. Right.

16 Q. Right. So much like the study by Stocks  
17 and his team, which you say you don't agree with --

18 A. Right.

19 Q. -- much like the study of Dr. Raval and  
20 his team, which you don't agree with; much like the  
21 statements of Mr. Olmstead, 3M's retained  
22 consultant, you don't agree with, you also don't  
23 agree with the seven researchers in this study that  
24 that can be used as a proxy between particulates  
25 and airborne biological matter?



1           A.    -- the interpretation of data is so  
2           important.

3           Q.    I'm not asking you if it's important.  
4           Why --

5           A.    Authors --

6           Q.    Why are they wrong?

7           A.    Authors always wish to say things that  
8           they set out to say. You know that.

9           Q.    Hold on. Are you -- are you claiming  
10          that Dr. Darouiche set out to prove a certain  
11          proposition in this report?

12          A.    Well, maybe --

13          Q.    What evidence do you have of that, sir?

14          A.    I'll let you in on a dirty little secret  
15          in that the purpose of somebody going to do a bunch  
16          of experiments is to hopefully get data to back up  
17          his expectation in the first place.

18          Q.    So you're saying --

19          A.    Are you surprised --

20          Q.    -- that you believe --

21          A.    -- to hear that?

22          Q.    I'm not -- I'm not surprised to hear  
23          anything from you today.

24          A.    Because --

25          Q.    What I'm -- what I'm asking you is do

1                                   You don't know Dr. Darouiche?

2                   A.    No.

3                   Q.    You don't know what his motivations were  
4                   in doing this study?

5                   A.    No.

6                   Q.    Do you know what his hypothesis was?

7                   A.    Well --

8                   Q.    Do you know?

9                   A.    He thinks that first and foremost he  
10                  could connect the infections that he saw with the  
11                  concentration of culturable particles in the air.

12                  Q.    That's his conclusion; right?

13                  A.    He set out to show that.

14                  Q.    Where do you see that he set out to do  
15                  that? Where is his hypothesis, sir?

16                  A.    Well, it's in the introduction.

17                  Q.    Where does it say that he wanted to  
18                  prove that this was true?

19                  A.    Well, if that's not what he wants to  
20                  show, then why bother to do any work?

21                  Q.    So we don't do science unless we have an  
22                  agenda? Is that what you're saying, sir?

23                  A.    Well, look at the title. The whole  
24                  title says that's what he intends to do.

25                  Q.    You think that the title --

1 A. Yeah.

2 Q. -- which represents the findings of this  
3 study --

4 A. Yeah.

5 Q. -- represents what his agenda was in  
6 doing this study?

7 A. Exactly.

8 MR. GORDON: Objection,  
9 argumentative.

10 MR. BANKSTON: All right.

11 Q. So according to you, Dr. Darouiche's  
12 work and the work of his entire team is tainted  
13 because apparently they had some sort of agenda or  
14 motivation?

15 A. I didn't say that. I simply say that  
16 the data that is presented does not support the  
17 statement they are making.

18 Q. Why not?

19 A. Well, as I've said, look at the --

20 Q. I'm looking at it. Tell me why.

21 A. Look at Figure 4 and 5 in particular.  
22 Look at the -- are you familiar with the 95 percent  
23 confident interval?

24 Q. Pretend I'm not.

25 A. Okay. Look at -- look at Y Figure 4.

1 Q. Okay. You knew that study is not  
2 controlled; right?

3 A. What do you mean by "controlled?"

4 Q. I mean, it has -- it's not a controlled  
5 study. You understand that?

6 A. You mean they did not use a control  
7 experiment?

8 Q. Correct.

9 A. Okay.

10 Q. Right. And you knew that from reading  
11 Dr. Yadin David's report; correct?

12 A. Yeah.

13 Q. That that Huang study was not  
14 controlled?

15 A. Right.

16 Q. All right. Now let me ask you something  
17 about controlled experiments. The Huang study is  
18 still useful to you; right? You still find it to  
19 be a useful study?

20 A. Come again.

21 Q. The Huang study --

22 A. Right.

23 Q. -- is a useful study to you? You find  
24 it to be a useful piece of literature?

25 A. Yeah, I would say that.

1 Q. Yeah. You cited it as support for some  
2 of your opinions you're giving; correct?

3 A. Right.

4 Q. Now, what I want to know is does the  
5 fact that Huang in his biological study, does the  
6 fact that he didn't do a controlled study, does  
7 that mean that he's not familiar with  
8 microbiological concepts?

9 A. I can't state that. I can't say that.

10 Q. Okay. So just because somebody doesn't  
11 use a control, that doesn't necessarily mean that  
12 they're not familiar with how to do a proper  
13 microbiological study? Is that your testimony?

14 A. That's fair.

15 Q. Okay. Thank you, sir. You know, again,  
16 and I'll just circle back because we're going to  
17 cover them again, Huang is one of these studies  
18 that involved a predecessor model Bair Hugger in  
19 2002; correct?

20 A. If you say so, yes.

21 Q. Well, you reviewed -- you're  
22 responding -- one of the things you're doing in  
23 this case is responding to the report of Dr. Yadin  
24 David; right?

25 A. You see, I -- when I was starting to

1 Q. Did you have any assistance in the  
2 writing of your report?

3 A. No.

4 Q. So this section here, you wrote this?

5 A. Yeah.

6 Q. Okay. Let's talk about page 18. Do you  
7 see where the first full paragraph talks about  
8 Albrecht and his colleagues?

9 A. Right.

10 Q. Okay. You talked -- you make one point,  
11 but what I want to go to is your second point here.  
12 Do you see where you say "On their second aim..."?

13 A. Right.

14 Q. Okay. It says: "On their second aim,  
15 it would appear [that] the authors were not  
16 familiar with microbiological concepts as the  
17 experimental design had no control." Correct?

18 A. Right.

19 Q. Do you remember when we talked about the  
20 Huang study and it not having a control?

21 A. You -- you mentioned that, yes.

22 Q. Yes. And you told me that the fact that  
23 it didn't have a control doesn't mean that they  
24 weren't unfamiliar with microbiological concepts.  
25 Do you remember telling me that?

1 A. I might have said that.

2 Q. Yeah. And you're saying the exact  
3 opposite here about Mr. Albrecht, aren't you?  
4 You're saying the fact that he didn't have a  
5 control means that he's unfamiliar with biological  
6 concepts. Correct?

7 A. Yes.

8 Q. So when it comes to literature that  
9 hurts 3M's case, you make what's essentially a  
10 criticism of this author. You attack his  
11 qualifications and his credibility saying he's not  
12 familiar with microbiological concepts because he  
13 had no control. But when you had a study that was  
14 favorable to the client who has hired you, you  
15 didn't mention that it wasn't controlled, nor did  
16 you criticise those authors, did you?

17 A. No.

18 Q. And you knew when you wrote your report  
19 that Huang was not controlled?

20 A. No.

21 Q. You did know that; correct?

22 A. Well, it might have -- I might have  
23 noted that, but it wasn't something that I jumped  
24 on.

25 Q. Right. Because you're only going to

1 insult an author if he's critical of 3M, not if  
2 he's in favor of 3M; right?

3 A. Right.

4 Q. Yeah. You're not going to criticise  
5 authors that are favorable of 3M; correct?

6 A. Right.

7 Q. Understood. So you remember back when  
8 we were talking about are you writing an objective  
9 independent report or are you doing a report to  
10 advocate for 3M, this is a report which is bias in  
11 favour of 3M, isn't it?

12 A. Are you saying that?

13 Q. I'm asking you if you believe it.  
14 Considering that you criticized one author for  
15 something, told us he wasn't familiar with  
16 microbiological concepts, but knowingly didn't even  
17 include the fact that this other author had no  
18 control and didn't criticise him at all, that's a  
19 form of bias, isn't it?

20 A. No.

21 Q. You don't think that's bias?

22 A. No.

23 Q. You don't think it's a little bit unfair  
24 to insult Mr. Albrecht and accuse him of having no  
25 familiarity of microbiological concepts --